

IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

Application No.: 10/669,824)
In re application of: JIANG, Cai-Zhong)
Filed: 23 September 2003)
Art Unit: 1638)
Examiner: KRUSE, David H.)
Docket No. MBI-0034CIP)
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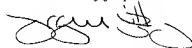
DECLARATION UNDER 37 CFR 1.132 OF JEFFREY M. LIBBY

I, Jeffrey M. Libby, declare:

1. I received my Bachelor of Science degree in Microbiology from the University of Illinois and my doctoral degree in Microbiology and Microbial Genetics from Cornell University. I joined Mendel Biotechnology in June 2002 and have served as Senior Patent Agent since June 2002. I state that I have prepared Exhibit B and Exhibit C for the response to the most recent Office action of the present patent application. I have determined the theoretical melting temperatures of G3456 and the homologous polynucleotide subsequences listed in Exhibit C by first aligning each pair of full length polynucleotide sequences, finding similar subsequences of length 50 bases within each aligned pair of sequences, and comparing the reverse complement of each listed homolog subsequence with the similar subsequence of G3456 using the DINAMelt server available at www.bioinfo.rpi.edu/applications/hybrid/hybrid2.php. This declaration is being drafted as part of my normal duties to support intellectual property at Mendel Biotechnology, Inc. As compensation for employment at Mendel Biotechnology, I receive salary, benefits and stock options.

2. I hereby declare that all statements made herein are true and that they are based on my own knowledge, information and belief. These statements are made with the knowledge that willful false statements are punishable by fine or imprisonment, or both, under Section 1001 of Title 18 of the United States Code and that such willful false statements may jeopardize the validity of this application or any patent issued from it.

Date: 7 January 2008



Jeffrey M. Libby, Ph.D.
Senior Patent Agent
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Exhibit B. Sequences phylogenetically-related to G3456, polypeptide SEQ ID NO: 14

The G3456 sequence used to generate the data in Table 1 of the attached declaration differs from the G3456 polypeptide sequence described in present specification by two residues, indicated below in small boxes.

>G3456 (showing residues of the polypeptide predicted to have been expressed in plants)
MKKPD~~L~~GF~~S~~SMNE~~S~~TV~~T~~GNHIGEEDE~~D~~RENSDEPREGAIDVATRRPRGRPPGSRNKP~~K~~PP
IFVTRDS~~P~~NALRSHVMEI~~A~~VGAD~~I~~AD~~C~~V~~A~~QFARRRQ~~R~~GV~~S~~ILSGSGTVVNVNLRQ~~T~~APG
AVMALHGRFDILSLTGSF~~L~~PGPSPPGATGLTIYLAGGQ~~G~~QIVG~~G~~QVVGPLV~~A~~AGPVLVMA
ATFSNATYERLPLEDDDDQEQHGSGGGGGSPQEK~~T~~GGPGEASSISIVYNNNNVPPSLGLP~~N~~G
QHLNHEAYSSPWGHS~~P~~SHARPPF*

The G3456 sequence described in the present specification is indicated below. The percentage identities of the second conserved domains of either the above or below sequence to the related sequences in Table 1 of the attached declaration remain the same regardless of which G3456 sequence is used to generate the comparisons.

>G3456 (Predicted polypeptide, 2nd conserved domain underlined)
MKKPD~~L~~GF~~S~~SMNE~~S~~TV~~T~~GNHIGEEDE~~D~~RENSDEPREGAIDVATRRPRGRPPGSRNKP~~K~~PP
IFVTRDS~~P~~NALRSHVMEI~~A~~VGAD~~I~~AD~~C~~V~~A~~QFARRRQ~~R~~GV~~S~~ILSGSGTVVNVNLRQ~~T~~APG
AVMALHGRFDILSLTGSF~~L~~PGPSPPGATGLTIYLAGGQ~~G~~QIVG~~G~~QVVGPLV~~A~~AGPVLVMA
ATFSNATYERLPLEDDDDQEQHGSGGGGGSPQEK~~T~~GGPGEASSISIVYNNNNVPPSLGLP~~N~~G
QHLNHEAYSSPWGHS~~P~~SHARPPF*

Related sequences in Table 1 of the attached declaration, their second conserved domains, and alignments to the G3456 second conserved domain are shown below.

>G3460 (Predicted polypeptide, 2nd conserved domain underlined)
MAGLDLGSASRFVQNLHLPLHLQ~~Q~~NYQQPRHKRDSEEQETPPNPGTALAPFDNDDK~~S~~Q
GLELASPGDIVGRRPRGRPSGSKNKP~~K~~PPVITRESANTLRAHILEVSGSGSDVFD~~C~~V~~T~~A
YARRRQ~~R~~GICVL~~S~~SGSGTVTNVSLRQPAAGAVVRLHGRFEILSLSGSF~~L~~PPPPAPP~~G~~ATSL
TIYLAGGQ~~G~~QVVG~~G~~NVVGELTAAGPVIVIAASFTNVAYERLPLEEDEQ~~Q~~Q~~Q~~QLIQSPAT
TSSQGNNNNNFPDPSSGLPFFNLPLNMQNVQLP~~P~~F*

Identity of G3456 second conserved domain to G3460 second conserved domain determined using manual alignment = 72/96 identical residues (75.0%)

G3456:	VAQFARRRQ R GV S ILSGSGTVVNVNLRQPTAPGAVMALHGRFDILSLTGSF L PGPSPPGA V +ARRRQ R G+ +LSGSGTV NV+LRQ P A GAV+ LHGRF+ILSL+GSFLP P+PPGA
G3460:	V T AYARRRQ R GICVL S SGSGTVTNVSLRQPAAGAVVRLHGRFEILSLSGSF L PPPPAPP G A
G3456:	TGLTIYLAGGQ G QIVGGEVVGPLV A AGPVLVMAATF T LTIYLAGGQ G Q+VGG VVG L AAGPV+V+AA+F
G3460:	TS L TIYLAGGQ G QVVG G NVVGELTAAGPVIVIAAS F

>G3459 (Predicted polypeptide, 2nd conserved domain underlined)
MAGLDLGLGSARFVNQLHRPDLHLQQNFQQHQDQHQRDLEEKTPPNHRMGAPFDDSD
RSPGLELTSGPGDIVGRRPRGRPPGSKNPKPPVITRESANTLRAHILEVSGSDVDFDC
VTAYARRRQRGICVLSGSGTVTNVSLRQPAAGAVVTLHGRFEILSLSGSFLPPAPPGA
TSLTIYLAGGQGVVGGNVIGELTAAGPVIVIAASFNTVAYERLPLEEDEQQQQQQLQI
QPPATTSSQGNNNNNNPFDPSSGLPFFNLPLMNQNVQLPVEGWAVNPASRPQPF*

Identity of G3456 second conserved domain to G3459 second conserved domain determined using
manual alignment = 71/96 identical residues (73.9%)

G3456:	VAQFARRRQRGVSI	LSGSGTVVNVNLRQPTAPGAVMALHGRFDILSLTGSFLPGSPPGA
	V	+ARRRQRG+ +LSGSGTV NV+LRQP A GAV+ LHGRF+ILSLT+GSFLP P+PPGA
G3459:	VTAYARRRQRGICVLSGSGTVTNVSLRQPAAGAVVTLHGRFEILSLSGSFLPPAPPGA	
G3456:	TGLTIYLAGGQGVVGGNVIGELTAAGPVIVIAASF	
	T	LTIIYLAGGQGV+VGG V+G L AAGPV+V+AA+P
G3459:	TSLTIYLAGGQGVVGGNVIGELTAAGPVIVIAASF	

>G2153 (Predicted polypeptide, 2nd conserved domain underlined)
MANPWWTGVNLSGLETPPGSSQLKKPDLHISMNMAMDSGHNNHHHHQEVNDDDD
DNLSDGDHPEPREGAVEAPTRRRPRGRPAGSKNPKPPIFVTRDSPNALKSHVMEIASGTV
IETLATFARRRQRGICILSGNGTVANVTLRQPSTAATAVAAAPGGAVALAQGRFEILSLTG
SFLPGPAPPGSTGLTIYLAGGQGVVGGSVVGPLMAAGPVMLIAATFNSNATYERLPLEEE
EAAERGGGGSGGVVPGQLGGGGPLSSGAGGGDGNQGLFPVYNMPPGNLVSNGSGGGGGG
SGQAYGWAQARSGF*

Identity of G3456 second conserved domain to G2153 second conserved domain determined using
manual alignment = 77/104 identical residues (74.0%)

G3456:	VAQFARRRQRGVSI	LSGSGTVVNVNLRQPT-----APG--AVMALHGRFDILSLTGSFL
	+A FARRRQRG+ ILSG+GTV NV LRQP+ APG AV+AL GRF+ILSLTGSFL	
G2153:	LATFARRRQRGICILSGNGTVANVTLRQPSTAATAVAAAPGGAVALAQGRFEILSLTGSFL	
G3456:	PGPSPPGATGLTIYLAGGQGVVGGSVVGPLVAAGPVLMVAATF	
	PGP+PPG+TGLTIYLAGGQGV+VGG VVGPL+AAGPV+++AATF	
G2153:	PGPAPPGSTGLTIYLAGGQGVVGGSVVGPLMAAGPVMLIAATF	

>G3401 (Predicted polypeptide, 2nd conserved domain underlined)
MASKEPSGDHDEMNGTSAGGGEPKDGAVVTVGNRRRPRGRPPGSKNPKPPIFVTRDSPN
ALRSHVMEVAGGADVAESIAHFARRRQRGVCVLSGAGTVTDVALRQPAAPSAVVALRGFR
EILSLTGTFLPGPAPPGSTGLTVYLAGGQGVVGGSVVGTTLTAAGPVMVIASTFANATYE
RLPLDQEEEEAAAGMMAPPPLMAGAADPLLFGGGMHDAGLAAWHHARP PPPPPY*

Identity of G3456 second conserved domain to G3401 second conserved domain determined using
manual alignment = 72/96 identical residues (75.0%)

G3456:	VAQFARRRQRGVSI	LSGSGTVVNVNLRQPTAPGAVMALHGRFDILSLTGSFLPGSPPGA
	+A FARRRQRGV +LSG+GTV +V LRQP AP AV+AL GRF+ILSLTG+FLPGP+PPG+	
G3401:	IAHFARRRQRGVCVLSGAGTVTDVALRQPAAPSAVVALRGFRFEILSLTGTFLPGPAPPGS	
G3456:	TGLTIYLAGGQGVVGGNVIGELTAAGPVIVIAASF	
	TGLT+YLAGGQGV+VGG VVG L AAGPV+V+A+TF	
G3401:	TGLTVYLAGGQGVVGGSVVGTTLTAAGPVMVIASTF	

>G3457 (Predicted polypeptide, 2nd conserved domain underlined)
MDPVAAQGRPLPPFFLTRDLHLHPHHQFQPHHNHQNTDEAGNGRGQKRDRDENAGGGGG
ATTPPQGGGEGKESGSGDGGGSDMGRPRGRPAGSKNKPKPPIIIITRDSANALRSHVMEI
ANGCDIMESITAFARRRQRGVCLSGSGTVTNVTLRQPASPGAVVTLHGRFEILSLSGSF
LPPPPAPPAASGLAIYLAGGQGQVVGGSVVGPLVASGPVVIMAASFGNAAYERLPLEEEET
PVAVAGNGGLGSPGIPGTQQQPQQQQQQQLVGDPNSSSLFHGMPQNLLNSVQLPAEGYWG
GSARPPF*

Identity of G3456 second conserved domain to G3457 second conserved domain determined using
manual alignment = 72/96 identical residues (75.0%)

G3456:	VAQFARRRQRGVSILSGSGTVVNVNLRQPTAPGAVMALHGRFDILSLTGSFLPGSPPGA
	+ FARRRQRGV +LSGSGTV NV LRQP +PGAV+ LHGRF+ILSL+GSFLP P+PP A
G3457:	ITAFARRRQRGVCLSGSGTVTNVTLRQPASPGAVVTLHGRFEILSLSGSFLEPPPPAPPA
G3456:	TGLTIYLAGGQGQIVGGEVVGGLVAAGPVLVMAATF
	+GL IYLAGGQGQ+VGG VVGPLVA+GPV++MAA+F
G3457:	SGLAIIYLAGGQGQVVGGSVVGPLVASGPVVIMAASF

Exhibit C. Polynucleotide subsequences used for hybridization analysis

The best identity match of 50-base subsequences from G3456 and homolog DNAs are listed below and were used for determining theoretical melting temperatures. The first sequence in each pair is the 50 base subsequence derived from G3456, and the second sequence of each pair is the reverse complement of the corresponding subsequence from each optimally-aligned G3456 homolog. The $T_m(\text{conc})$ (the point at which the concentration of double-stranded molecules of one-half of its maximum value defines the melting temperature) was used to determine theoretical melting temperatures at 0.2x SSC (about 30 mM Na⁺) and 2.0x SSC (about 300 mM Na⁺). Determinations made with DINAMelt server available at: www.bioinfo.rpi.edu/applications/hybrid/hybrid2.php.

G3456 CCCTGGGCCGTCCCCTCCCGGCCACCGGGCTCACAATCTACCTCGCCG
G3456 CGGCGAGGTAGATTGTGAGCCCGGTGGCGCCGGGAGGGGACGGCCAGGG
 $T_m(\text{conc})$ 0.2x SSC: 82.3° C
 $T_m(\text{conc})$ 2x SSC: 93.6° C

G3456 GCCACCGGGCTCACAATCTACCTCGCCGAGGCCAGGGGCAGATCGTCGG
G3460 CCGACGACCTGGCCCTGCCCGCCGCGAGGTAGATTGTGAGACTGGTGCG
 $T_m(\text{conc})$ 0.2x SSC: 72.0° C
 $T_m(\text{conc})$ 2x SSC: 83.3° C

G3456 GCCACCGGGCTCACAATCTACCTCGCCGAGGCCAGGGGCAGATCGTCGG
G3459 CCGACAACCTGCCCTGCCCGCCGCGAGGTAGATTGTGAGGCTGGTGCG
 $T_m(\text{conc})$ 0.2x SSC: 71.8° C
 $T_m(\text{conc})$ 2x SSC: 80.9° C

G3456 GTCACCCGAGACAGCCCTAACGCGCTGCGGAGCCACGTCATGGAGATTGC
G2153 GCGATCTCCATGACATGGCTCTTGAGAGCATTGGAGAAATCGCGAGTGAC
 $T_m(\text{conc})$ 0.2x SSC: 70.0° C
 $T_m(\text{conc})$ 2x SSC: 77.3° C

G3456 CCTCCCGGCCGCCACCGGGCTCACAATCTACCTCGCCGGAGGGCAGGGGCA
G3401 TGCCCCCTGCCCGCCGCGAGGTACACGGTCAGCCCGGTGGAGCCCGGCGG
 $T_m(\text{conc})$ 0.2x SSC: 77.2° C
 $T_m(\text{conc})$ 2x SSC: 86.0° C

G3456 GTGGTGGGCCCACTCGTGGCGGGGGCCCCGTATTGGTAATGGCGGCTAC
G1073 GAAGCAGCCATCAACACTACCGGTCCCGAAGCAATTACGAACCGACAC
 $T_m(\text{conc})$ 0.2x SSC: 70.3° C
 $T_m(\text{conc})$ 2x SSC: 78.7° C